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| PPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. | |
|------------------------|-------------------------------|----------------------|---------------------|------------------|--|
| 10/763,854 | 01/22/2004 | Mark David Fidock | PC10959B | PC10959B 8327 | |
| 28523 | 7590 07/02/2004 | | EXAMINER | | |
| PFIZER INC | | LI, RUIXIANG | | | |
| PATENT DE EASTERN P | PARTMENT, MS8260 OINT ROAD | ART UNIT | PAPER NUMBER | | |
| GROTON, C | | 1646 | | | |
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DATE MAILED: 07/02/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

| | | Applicati | on No. | Applicant(s) | | | |
|--|--|-----------------------|---|--------------------|--------|--|--|
| Office Action Summary | | 10/763,8 | 54 | FIDOCK, MARK DAVID | | | |
| | | Examine | r | Art Unit | | | |
| | | Ruixiang | | 1646 | | | |
| The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply | | | | | | | |
| A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). | | | | | | | |
| Status | | | | | | | |
| 1) | Responsive to communication(s) file | ed on | | | | | |
| 2a)□ | This action is FINAL . | 2b)⊠ This action is r | non-final. | | | | |
| | the second secon | | | | | | |
| Disposition of Claims | | | | | | | |
| 4) Claim(s) 1-21 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration. 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-21 are subject to restriction and/or election requirement. | | | | | | | |
| Applicati | on Papers | | | | | | |
| 9)☐ The specification is objected to by the Examiner. | | | | | | | |
| 10) The drawing(s) filed on is/are: a) accepted or b) dojected to by the Examiner. | | | | | | | |
| Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). | | | | | | | |
| Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d). 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152. | | | | | | | |
| Priority under 35 U.S.C. § 119 | | | | | | | |
| 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. | | | | | | | |
| | | | | | | | |
| Attachment(s) | | | | | | | |
| | e of References Cited (PTO-892) | OTO-048) | 4) Interview Summary Paper No(s)/Mail D | | | | |
| 3) Inform | e of Draftsperson's Patent Drawing Review (F mation Disclosure Statement(s) (PTO-1449 or r No(s)/Mail Date | | 5) Notice of Informal F 6) Other: | | O-152) | | |

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Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - I. Claims 1-8, 10, and 19, drawn to an isolated polynucleotide, a vector, a host cell, and a process of producing a polypeptide, classified in class 536, subclass 23.5 and class 435, subclass 320.1, 325, and 69.1.
 - II. Claims 9, 11, and 12, drawn to an isolated polypeptide, classified in class 530, subclass 350.
 - III. Claims 13 and 14, drawn to an antibody, classified in class 530, subclass 387.9.
 - IV. Claims 15-18, drawn to a method for identifying a compound that binds and activate the polypeptide of claim 11 or inhibits the activation of the polypeptide of claim 11, classified in class 435, subclasses 6 and 7.1.
 - V. Claim 20, drawn to a method of elucidating the 3-dimensional structure of the polypeptide of claim 11, classified in class 530, subclass 412.
 - VI. Claim 21, drawn to a method of modeling the structure of the polypeptide of claim 11, classified in class 436, subclass 250.
- 2. The inventions are distinct, each from the other for the following reasons. Inventions I-III are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP §806.04, MPEP §808.01). In the instant case, the different inventions are drawn to completely different products, nucleic acid molecules, polypeptides, and antibodies. These molecules have completely different

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structures and biological functions which are not interchangeable and which require non-cohesive searches and considerations.

- 3. Inventions IV-VI are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP §806.04, MPEP §808.01). In the instant case, the different inventions are drawn to completely different methods each having completely different method steps, using different compositions, and having completely different outcomes. Invention IV is drawn to a method for identifying a compound that binds and activate the polypeptide of claim 11 or inhibits the activation of the polypeptide of claim 11, which is not required in Inventions V and VI; Invention V requires crystallizing a polypeptide and elucidating the structure of the polypeptide by X-ray crystallography, which is not required in Invention IV and VI; Invention Vi requires a method of modeling the polypeptide structure, which is not required in Invention IV and V. Thus, each method is unique and not required one for another and requires non-cohesive searches and considerations.
- 4. Inventions I and IV are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP §806.05 (h)). In the instant case, the nucleic acids may be used in a materially different process such as production of a polypeptide.

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- 5. Invention II is related to Inventions IV-VI as product and process of use. The invention can be shown to be distinct if either or both of the following can be shown:
 (1) the process for using the product as can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP §806.05 (h)). In the instant case, the polypeptide may be used in a materially different process such as to immunize mice to produce an antibody
- 6. Invention III is related to Invention IV as product and process of use. The invention can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP §806.05 (h)). In the instant case, the antibody may be used in a materially different process such as to immunoprecipitate or purify a polypeptide.
- 7. Invention I is an independent invention from Inventions V and VI; Invention III is an independent invention from Inventions V and VI. The different inventions are drawn to distinct product and method inventions.
- 8. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.
- 9. Because these inventions are distinct for the reasons given above and the search required for a single group is not required for any other group, restriction for examination purposes as indicated is proper.

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10. The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai, In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product

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claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48 (b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48 (b) and by the fee required under 37 CFR 1.17 (l).

Advisory Information

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ruixiang Li whose telephone number is (571) 272-0875. The examiner can normally be reached on Monday through Friday from 8:30 am to 5:00 pm. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Kunz, can be reached on (571) 272-0887.

Communications via Internet e-mail regarding this application, other than those under 35 U.S.C. 132 or which otherwise require a signature, may be used by the applicant and should be addressed to [yvonne.eyler@uspto.gov]. All Internet e-mail communications will be made of record in the application file. PTO employees do not engage in Internet communications where there exists a possibility that sensitive

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information could be identified or exchanged unless the record includes a properly signed express waiver of the confidentiality requirements of 35 U.S.C. 122. This is more clearly set forth in the Interim Internet Usage Policy published in the Official Gazette of the Patent and Trademark on February 25, 1997 at 1195 OG 89.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (571) 272-1600.

Ruixiang Li, Ph.D.

Examiner

June 28, 2004